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INFORMATION DISCLOSURE STATEMENT BY APPLICANT (use as many sheets as necessary)		Application Number	09/785,548
		Filing Date	2/20/2001
		First Named Inventor	Koutnikova, Hana
		Art Unit	1647
		Examiner Name	Hayes, Robert Clinton
Sheet 1 of 2	Attorney Docket Number	ST00005	

U.S. PATENT DOCUMENTS					
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FOREIGN PATENT DOCUMENTS						
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Examiner Signature	<i>R Hayes</i>	Date Considered	2/19/03
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Substitute for form 1449B/PTO		Complete if Known	
INFORMATION DISCLOSURE STATEMENT BY APPLICANT (use as many sheets as necessary)		Application Number	09/785,548
		Filing Date	2/20/2001
		First Named Inventor	Koutnikova, Hana
		Group Art Unit	1647
		Examiner Name	Hayes, Robert Clinton
		Attorney Docket Number	ST00005
Sheet	2	of	2

OTHER PRIOR ART -- NON PATENT LITERATURE DOCUMENTS

Examiner Initials	Cite No. 1	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published	T ²
no pay com pay w/SEPT w/SEPT	no pay com pay w/SEPT w/SEPT	STRAUSBERG, R.; Database Embl Sequences 'en ligne! ,Accession No. A1701008, "EST; H. sapiens cDNA clone IMAGE: 2340603 similar to TR: Q14184 DOC2-beta"; 6/4/1999.	
PKH		KITADA, T., et al; Mutations in the parkin gene cause autosomal recessive juvenile parkinsonism; Nature, Vol. 329, April 9,1998, 605-608.	
		ABBAS, N., et al.; A wide variety of mutations in the parkin gene are responsible for autosomal recessive parkinsonism in Europe; Hum. Mol. Genet., Vol. 8, 1999, 567-574.	
		HATTORI, N., et al.; Molecular genetic analysis of a novel Parkin gene in Japanese families with autosomal recessive juvenile parkinsonism; Ann Neurol., Vol. 6, 1998, 935-941.	
		LUCKING, C., et al.; Homozygous deletions in parkin gene in European and North African families with autosomal recessive juvenile parkinsonism; Lancet, Vol. 352, 1998, 1355-1356.	
		MORETT, E.; A novel transactivation domain in parkin; Trends Biochem Sci., Vol. 24, 1999, 229-231.	
		POLYMERPOULOS, M.H., et al.; Mutation in the alpha-synuclein gene identified in families with Parkinson's disease; Science, Vol. 276, 1997, 2045-2047.	
		SHIMURA, H., et al.; Immunohistochemical and subcellular localization of Parkin protein: absence of protein in autosomal recessive juvenile parkinsonism patients; Ann Neurol., Vol. 45, 1999, 668-672.	
		SUNADA, Y., et al.; Differential expression of the parkin gene in the human brain and peripheral leukocytes; Neurosci Lett., Vol. 254, 1998, 180-182.	
		MELLICK, G.D. et al.; The parkin gene S/N167 polymorphism in Australian Parkinson's disease patients and controls; Parkinsonism and Related Disorders 7, 2001, 89-91,	

Examiner Signature	<i>[Signature]</i>	Date Considered	2/14/03
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